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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/580,232	02/07/2007	Per Mansson	MANS3014/REF	3755
23364 BACON & TH	7590 02/18/201 OMAS, PLLC	EXAMINER		
625 SLATERS LANE			LUM, LEON YUN BON	
FOURTH FLOOR ALEXANDRIA, VA 22314-1176			ART UNIT	PAPER NUMBER
			1641	
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# Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)				
Office Action Comments	10/580,232	MANSSON ET AL.				
Office Action Summary	Examiner	Art Unit				
	Leon Y. Lum	1641				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply						
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status						
1) Responsive to communication(s) filed on <u>04 D</u>	ecember 2009					
,	·					
<i>i</i> —	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.					
closed in accordance with the practice under Lx parte Quayle, 1933 C.D. 11, 403 C.G. 213.						
Disposition of Claims						
4)⊠ Claim(s) <u>1-6,16 and 25-34</u> is/are pending in th	4)⊠ Claim(s) <u>1-6,16 and 25-34</u> is/are pending in the application.					
4a) Of the above claim(s) is/are withdra	4a) Of the above claim(s) is/are withdrawn from consideration.					
5) Claim(s) is/are allowed.						
6)⊠ Claim(s) <u>1-6,16 and 25-34</u> is/are rejected.						
7) Claim(s) is/are objected to.						
8) Claim(s) are subject to restriction and/o	r election requirement.					
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Application Papers						
9)☐ The specification is objected to by the Examiner.						
10)⊠ The drawing(s) filed on <u>22 May 2006</u> is/are: a)⊠ accepted or b)□ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the priority documents have been received in this National Stage						
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of the certified copies not received.						
Attachment(s)						
1) Notice of References Cited (PTO-892)  4) Interview Summary (PTO-413)						
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	Paper No(s)/Mail Da					
Information Disclosure Statement(s) (PTO/SB/08)     Paper No(s)/Mail Date	5) Notice of Informal P 6) Other:	atent Application				

### **DETAILED ACTION**

### Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* **v.** *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

- 1. Determining the scope and contents of the prior art.
- 2. Ascertaining the differences between the prior art and the claims at issue.
- 3. Resolving the level of ordinary skill in the pertinent art.
- 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

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Claims 1-5 and 25-32 are rejected under 35 U.S.C. 103(a) as being unpatentable over WO 00/43774 to Willner, cited in the IDS filed May 22, 2006, in view of U.S. Patent No. 5,047,326 to Pronovost.

## i. Independent claims 1 and 26 are obvious

Willner describes a competition immunoassay in which a sample comprising an analyte is contacted with a neutralizing agent, e.g. an antibody, and the mixture then contacted with a piezoelectric crystal with the assayed antigen thereon. See page 10, line 10 to page 11, line 7. With this description, Willner teaches an unlabeled antibody that is capable of binding to a target antigen for use in a piezoelectric crystal detection device. Willner, however, does not teach a mixture of at least two different unlabeled antibodies.

Pronovost describes a competitive immunoassay that utilizes a mixture of different antibodies, in order to detect the presence of multiple antigens corresponding to different biological entities. *See* column 4, lines 25-37.

With the foregoing description in mind, one of ordinary skill in the art would have found it obvious to modify Willner's antibody solution by including a mixture of different antibodies. The skilled artisan would have been motivated to make the modification because Provonost indicates that such a mixture of antibodies can detect multiple antigens. The skilled artisan would recognize that this mixture would therefore allow multiplexed detection (i.e., simultaneous detection of different antigen). Moreover, the skilled artisan would have had a reasonable expectation of success in combining the teachings of Willner and Provonost because it would have taken only routine skill in the

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art to immobilize different antigens on the same piezoelectric crystal in order to make use of Brooker's multiple antibodies to perform a multiplex competition immunoassay.

ii. Dependent claims 2- 5, 25 and 27-32 are obvious

Regarding claim 2, Willner and Provonost describe an antigen-antibody interaction pair. See supra rejection of claim 1.

Regarding claim 3, Willner teaches monoclonal antibodies. See page 8, line 3.

Regarding claims 4 and 28, Willner and Provonost do not explicitly describe an antibody concentration in the claimed range. Willner does, however, describe a displacement immunoassay method that utilizes a 0.1 mg/ml concentration of antibody. See page 29, lines 15-16. It would have been obvious to one of ordinary skill in the art to modify Willner and Provonost's antibody mixture, taught from the perspective of a competitive immunoassay, to limit the concentration of each antibody to between 0.1 and 0.8 mg/ml. Indeed, the skilled artisan would have arrived at the claimed range based on the doctrine of routine optimization. In a case decided by the precursor to the Federal Circuit, the court stated that a claim is not allowable where the skilled artisan could have arrived at the claim through routine experimentation on the optimum or workable ranges of the claim. In re Aller, 220 F.2d 454, 456 (CCPA 1955) (stating "where the general conditions of a claim are disclosed in the prior art, it is not inventive to discover the optimum or workable ranges by routine experimentation.") In Aller, the claims were directed to a process taught by the prior art, except for a specific temperature and acid concentration range. *Id*. The court, however, held that the claims were not patentable since the skilled artisan could have arrived at the claimed ranges through routine optimization.

The facts of *Aller* are relevant here. Similar to that case, Willner and Provonost teach all the limitations of claims 4 and 28, except for the specific concentration range. Willner does, however, teach an antibody concentration within the range, albeit in a different context (displacement versus competition immunoassay). But because the displacement and competition methods are described as alternative embodiments of the same method, the skilled artisan would have found it obvious to conduct routine experimentation to use the concentration described for the displacement assay and apply it to the competition assay.

Regarding claims 5, 27 and 32, Willner describes antibodies diluted in PBS. See page 21, line 8.

Regarding claims 25 and 31, Willner teaches DNT and TNT explosives. *See* page 5, line 8.

Regarding claims 29 and 30, Willner describes a probe solution that contains antibody at a fixed concentration. See page 32, lines 20-21.

Claims 16 and 33-34 are rejected under 35 U.S.C. 103(a) as being unpatentable over Willner in view of Provonost as applied to claim 1 above, and further in view of U.S. Patent No. 5,420,016 to Boguslaski *et al.* ("Boguslaski").

Willner and Provonost, described above, do not teach packaging the mixture into a kit.

Boguslaski teaches that assembling various assay system components into a test kit facilitates a convenient and facile use of the components. *See* column 7, lines 8-17.

With the foregoing description in mind, one of ordinary skill in the art would have found it obvious to modify Willner and Provonost's mixture by placing it in a kit. The skilled artisan would have made the modification because Boguslaski indicates that packaging components in a kit facilitates a convenient and facile use of the components. Moreover, because placing components in a kit involves mere routine skill in the art, the skilled artisan would have had a reasonable expectation of success.

Regarding claim 33, Willner teaches DNT and TNT explosives. *See* page 5, line 8.

Regarding claim 34, Provonost describes bottles and test tubes that can be used in conjunction with the competitive immunoassay described. *See* column 5, lines 24-26. It would have been obvious to one of ordinary skill in the art to use one of the bottles and/or test tubes to hold the antibody mixture prior to performing the immunoassay. Indeed, the mixture needs to be placed in such a container and the skilled artisan would recognize that a bottle or test tube would be suitable containers to store the mixture.

Claim 6 is rejected under 35 U.S.C. 103(a) as being unpatentable over Willner and Provonost as applied to claim 1 above, and further in view of U.S. Patent No. 4,375,414 to Strahilevitz.

Willner and Provonost, described above, do not teach a narcotic analyte.

Strahilevitz describes an antibody directed to heroin, in order to detect the drug in a biological material. See abstract; column 1, lines 13-15.

With the foregoing description in mind, one of ordinary skill in the art would have found it obvious to modify Willner and Provonost's antibody mixture to include an antibody to heroin. This modification would allow a user to detect heroin in a biological material, thereby providing a reason for the skilled artisan to perform the modification. Moreover, because the modification simply substitutes one antibody for another in a method that can be applied generally to an antibody, the skilled artisan would have had a reasonable expectation of success.

### Response to Arguments

Applicant's arguments, see Response filed December 4, 2009, with respect to the rejection of independent claims 1 and 26 under 35 U.S.C. 103(a) have been fully considered and are persuasive. Therefore, the rejection has been withdrawn. However, upon further consideration, a new ground of rejection is made in view of Willner and Provonost. *See supra* rejection of claims 1 and 26.

#### Conclusion

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Leon Y. Lum whose telephone number is (571) 272-

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2872. The examiner can normally be reached on Monday to Friday (8:30 am to 5:00

pm).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Mark L. Shibuya can be reached on (571) 272-0806. The fax phone number

for the organization where this application or proceeding is assigned is 571-273-8300.

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/Leon Y. Lum/

Examiner, Art Unit 1641

/Unsu Jung/

Primary Examiner, Art Unit 1641